12 December 2019
EMA/270645/2015
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: oxaliplatin

Procedure no.: PSUSA/00002229/201804

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Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Axiplatin 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	88849.00.00	AXIOS PHARMA GMBH	DE
axiplatin 5 mg/ml Pulver zur Herstellung einer Infusionslösung	not available	75516.00.00	APOCARE PHARMA GMBH	DE
Bendaplatin 5 mg/ml Pulver zur Herstellung einer Infusionslösung	DE/H/2980/001	68364.00.00	BENDALIS GMBH	DE
Croloxat® 5 mg/ml Pulver zur Herstellung einer Infusionslösung	FI/H/0589/001	66245.00.00	STADAPHARM GMBH	DE
DACPLAT 5 mg/ml, poudre pour solution pour perfusion	not available	34009 563 917 8 3	MEDIPHA SANTE	FR
DACPLAT 5 mg/ml, poudre pour solution pour perfusion	not available	34009 559 645 7 5	MEDIPHA SANTE	FR
DACPLAT 5 mg/ml, poudre pour solution pour perfusion	not available	34009 559 646 3 6	MEDIPHA SANTE	FR
DACPLAT 5 mg/ml, solution à diluer pour perfusion	FR/H/0285/002	34009 567 123 6 6	MEDIPHA SANTE	FR
DACPLAT 5 mg/ml, solution à diluer pour perfusion	FR/H/0285/002	34009 567 124 2 7	MEDIPHA SANTE	FR
DACPLAT 5 mg/ml, solution à diluer pour perfusion	FR/H/0285/002	34009 569 816 9 4	MEDIPHA SANTE	FR
Elatofen 5 mg / ml, σκόνη για διάλυμα προς έγχυση	EL/H/0162/002	31490/11/9-1-12	DEMO ABEE	GR
Elatofen 5 mg / ml, σκόνη για διάλυμα προς έγχυση	EL/H/0162/002	79963/12-11-2012	DEMO ABEE	GR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ELOXATIN	not available	20296	SANOFI-AVENTIS CYPRUS LTD	CY
ELOXATIN	not available	20296	SANOFI-AVENTIS CYPRUS LTD	CY
ELOXATIN	not available	20296	SANOFI-AVENTIS CYPRUS LTD	CY
ELOXATIN	FR/H/0284/002	38482	SANOFI-AVENTIS DENMARK A/S	DK
ELOXATIN	FR/H/0284/002	38482	SANOFI-AVENTIS DENMARK A/S	DK
ELOXATIN	FR/H/0284/002	38482	SANOFI-AVENTIS DENMARK A/S	DK
ELOXATIN	FR/H/0284/002	269890201	SANOFI-AVENTIS AEBE	GR
ELOXATIN	FR/H/0284/002	269890203	SANOFI-AVENTIS AEBE	GR
ELOXATIN	FR/H/0284/002	269890202	SANOFI-AVENTIS AEBE	GR
ELOXATIN	FR/H/0284/002	MA 082/00502	SANOFI MALTA LTD	MT
ELOXATIN	FR/H/0284/002	MA 082/00502	SANOFI MALTA LTD	MT
ELOXATIN	FR/H/0284/002	MA 082/00502	SANOFI MALTA LTD	MT
ELOXATIN 5 MG/ML	not available	5600/2013/01	SANOFI ROMANIA SRL	RO
ELOXATIN 5 MG/ML	not available	5600/2013/02	SANOFI ROMANIA SRL	RO
ELOXATIN 5 MG/ML	not available	5600/2013/03	SANOFI ROMANIA SRL	RO
ELOXATIN 5 MG/ML CONCENTRAAT	FR/H/0144/002	RVG 32774	SANOFI-AVENTIS NETHERLANDS B.V.	NL
ELOXATIN 5 MG/ML CONCENTRAAT	FR/H/0144/002	RVG 32774	SANOFI-AVENTIS NETHERLANDS B.V.	NL
ELOXATIN 5 MG/ML CONCENTRAAT	FR/H/0144/002	RVG 32774	SANOFI-AVENTIS NETHERLANDS B.V.	NL
ELOXATIN 5 mg/ml concentraat voor oplossing voor infusie	FR/H/0144/002	BE281601	SANOFI BELGIUM	BE
ELOXATIN 5 mg/ml concentraat voor oplossing voor infusie	FR/H/0144/002	BE281592	SANOFI BELGIUM	BE
ELOXATIN 5 mg/ml	FR/H/0144/002	BE288784	SANOFI BELGIUM	BE

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concentraat voor oplossing voor infusie				
Eloxatin 5 mg/ml concentrado para solución para perfusión.	FR/H/0144/002	67390	SANOFI-AVENTIS, S.A.	ES
Eloxatin 5 mg/ml concentrado para solución para perfusión.	FR/H/0144/002	67390	SANOFI-AVENTIS, S.A.	ES
Eloxatin 5 mg/ml concentrado para solución para perfusión.	FR/H/0144/002	67390	SANOFI-AVENTIS, S.A.	ES
ELOXATIN 5 MG/ML CONCENTRATE FOR SOLUTION FOR INFUSION SOLUTION	not available	20060432	SANOFI BULGARIA EOOD	BG
ELOXATIN 5 MG/ML CONCENTRATE FOR SOLUTION FOR INFUSION SOLUTION	not available	20060432	SANOFI BULGARIA EOOD	BG
ELOXATIN 5 MG/ML CONCENTRATE FOR SOLUTION FOR INFUSION SOLUTION	not available	20060432	SANOFI BULGARIA EOOD	BG
ELOXATIN 5 mg/ml concentrato per soluzione per infusione	FR/H/0144/002	034411037	SANOFI S.P.A	IT
ELOXATIN 5 mg/ml concentrato per soluzione per infusione	FR/H/0144/002	034411052	SANOFI S.P.A	IT
ELOXATIN 5 mg/ml concentrato per soluzione per infusione	FR/H/0144/002	034411049	SANOFI S.P.A	IT
Eloxatin 5 mg/ml koncentrat za otopinu za infuziju	not available	UP/I-530-09/13-02/54	SANOFI-AVENTIS CROATIA D.O.O.	HR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ELOXATIN 5 mg/ml koncentrat za raztopino za infundiranje	FR/H/0284/002	H/06/00540/002	SANOFI-AVENTIS D.O.O.	SI
ELOXATIN 5 mg/ml koncentrat za raztopino za infundiranje	FR/H/0284/002	H/06/00540/001	SANOFI-AVENTIS D.O.O.	SI
ELOXATIN 5 mg/ml koncentrat za raztopino za infundiranje	FR/H/0284/002	H/06/00540/003	SANOFI-AVENTIS D.O.O.	SI
ELOXATIN 5 MG/ML KONZENTRAT ZUR HERSTELLUNG EINER INFUSIONSLOESUNG	FR/H/0284/002	1-26251	SANOFI-AVENTIS GMBH OSTERREICH	AT
ELOXATIN 5 MG/ML KONZENTRAT ZUR HERSTELLUNG EINER INFUSIONSLOESUNG	FR/H/0284/002	1-26251	SANOFI-AVENTIS GMBH OSTERREICH	AT
ELOXATIN 5 MG/ML KONZENTRAT ZUR HERSTELLUNG EINER INFUSIONSLOESUNG	FR/H/0284/002	1-26251	SANOFI-AVENTIS GMBH OSTERREICH	AT
ELOXATIN 5 MG/ML KONZENTRAT ZUR HERSTELLUNG EINER INFUSIONSLOESUNG	FR/H/0144/002	63264.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
ELOXATIN 5 MG/ML KONZENTRAT ZUR HERSTELLUNG EINER INFUSIONSLOESUNG	FR/H/0144/002	63264.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
ELOXATIN 5 MG/ML KONZENTRAT ZUR HERSTELLUNG EINER INFUSIONSLOESUNG	FR/H/0144/002	63264.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
ELOXATIN 5 mg/ml polvere per soluzione per	FR/H/0144/001	034411013	SANOFI S.P.A	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
infusione				
ELOXATIN 5 mg/ml polvere per soluzione per infusione	FR/H/0144/001	034411025	SANOFI S.P.A	IT
ELOXATIN 5 MG/ML POWDER FOR SOLUTION FOR INFUSION	not available	019040	SANOFI-AVENTIS CYPRUS LTD	CY
ELOXATIN 5 MG/ML POWDER FOR SOLUTION FOR INFUSION	not available	019039	SANOFI-AVENTIS CYPRUS LTD	CY
ELOXATIN 5 mg/ml solution à diluer pour perfusion	FR/H/0144/002	BE281592	SANOFI BELGIUM	BE
ELOXATIN 5 mg/ml solution à diluer pour perfusion	FR/H/0144/002	BE281601	SANOFI BELGIUM	BE
ELOXATIN 5 mg/ml solution à diluer pour perfusion	FR/H/0144/002	BE288784	SANOFI BELGIUM	BE
ELOXATIN 5 mg/ml solution à diluer pour perfusion	FR/H/0144/002	0422335	SANOFI BELGIUM	LU
ELOXATIN 5 mg/ml solution à diluer pour perfusion	FR/H/0144/002	0436599	SANOFI BELGIUM	LU
ELOXATIN 5 mg/ml solution à diluer pour perfusion	FR/H/0144/002	0422321	SANOFI BELGIUM	LU
Eloxatin 5 mg/ml concentrate for solution for infusion	FR/H/0284/002	PA 540/148/1	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Eloxatin 5 mg/ml concentrate for solution for infusion	FR/H/0284/002	PA 540/148/1	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Eloxatin 5 mg/ml	FR/H/0284/002	PA 540/148/1	SANOFI-AVENTIS IRELAND	IE

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concentrate for solution for infusion			LTD. T/A SANOFI	
Eloxatin, 5 mg/ml, concentrado para solução para perfusão	FR/H/0284/002	5865886	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Eloxatin, 5 mg/ml, concentrado para solução para perfusão	FR/H/0284/002	5710082	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Eloxatin, 5 mg/ml, concentrado para solução para perfusão	FR/H/0284/002	5710181	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
ELOXATINE 5 MG/ML CONCENTRATE FOR SOLUTION FOR INFUSION	FR/H/0144/002	PL 04425/0296	AVENTIS PHARMA LTD	UK
ELOXATINE 5 MG/ML CONCENTRATE FOR SOLUTION FOR INFUSION	FR/H/0144/002	PL 04425/0296	AVENTIS PHARMA LTD	UK
ELOXATINE 5 MG/ML CONCENTRATE FOR SOLUTION FOR INFUSION	FR/H/0144/002	PL 04425/0296	AVENTIS PHARMA LTD	UK
ELOXATINE 5 MG/ML, POUDRE POUR SOLUTION POUR PERFUSION	FR/H/0144/001	559 649-2	SANOFI-AVENTIS FRANCE	FR
ELOXATINE 5 MG/ML, POUDRE POUR SOLUTION POUR PERFUSION	FR/H/0144/001	559 648-6	SANOFI-AVENTIS FRANCE	FR
ELOXATINE 5 MG/ML, POUDRE POUR SOLUTION POUR PERFUSION	FR/H/0144/001	563 191-7	SANOFI-AVENTIS FRANCE	FR

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ELOXATINE 5 MG/ML, SOLUTION A DILUER POUR PERFUSION	FR/H/0144/002	34009 565 984 4 1	SANOFI-AVENTIS FRANCE	FR
ELOXATINE 5 MG/ML, SOLUTION A DILUER POUR PERFUSION	FR/H/0144/002	34009 565 983 8 0	SANOFI-AVENTIS FRANCE	FR
ELOXATINE 5 MG/ML, SOLUTION A DILUER POUR PERFUSION	FR/H/0144/002	34009 569 560 4 3	SANOFI-AVENTIS FRANCE	FR
EUROXALIPLATIN 5 mg/ml	not available	79591.00.00	LAPHARM GMBH PHARMAZEUTISCHE PRODUKTE	DE
Euroxaliplatin 5 mg/ml Konzentrat	not available	89390.00.00	LAPHARM GMBH PHARMAZEUTISCHE PRODUKTE	DE
GENEPLATIN® 5mg/ml πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση	not available	95296	GENEPHARM S.A.	GR
Gessedil 5 mg/ml por oldatos infúzióhoz	not available	OGYI-T-21080/01-02	EGIS PHARMACEUTICALS PLC	HU
Medoxa 5 mg/ml Pulver zur Herstellung einer Infusionslösung	FI/H/0584/001	66241.00.00	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	DE
Oksaliplatin Kabi 5 mg/ml koncentrat za raztopino za infundiranje	AT/H/0893/001	H/11/01147/001	FRESENIUS KABI ONCOLOGY PLC.	SI
Oksaliplatin Kabi 5 mg/ml koncentrat za raztopino za infundiranje	AT/H/0893/001	H/11/01147/002	FRESENIUS KABI ONCOLOGY PLC.	SI
Oksaliplatin Kabi 5 mg/ml koncentrat za raztopino za infundiranje	AT/H/0893/001	H/11/01147/003	FRESENIUS KABI ONCOLOGY PLC.	SI
Oksaliplatin Mylan 5	NL/H/1977/001	H/12/01149/001	MYLAN S.A.S	SI

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mg/ml koncentrat za raztopino za infundiranje				
Oksaliplatin Mylan 5 mg/ml koncentrat za raztopino za infundiranje	NL/H/1977/001	H/12/01149/002	MYLAN S.A.S	SI
Oksaliplatin Mylan 5 mg/ml koncentrat za raztopino za infundiranje	NL/H/1977/001	H/12/01149/003	MYLAN S.A.S	SI
Oksaliplatin SUN 5 mg/ml konsentrat til infusjonsvæske, oppløsning	UK/H/4547/001	12-8909	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	NO
Oksaliplatin Teva 5 mg/ml koncentrat za raztopino za infundiranje	NL/H/0820/001	H/07/01150/003	TEVA PHARMA B.V.	SI
Oksaliplatin Teva 5 mg/ml koncentrat za raztopino za infundiranje	NL/H/0820/001	H/07/01150/001	TEVA PHARMA B.V.	SI
Oksaliplatin Teva 5 mg/ml koncentrat za raztopino za infundiranje	NL/H/0820/001	H/07/01150/002	TEVA PHARMA B.V.	SI
Oksaliplatin Teva 5 mg/ml koncentrat za raztopino za infundiranje	NL/H/0820/001	H/07/01150/004	TEVA PHARMA B.V.	SI
Oksaliplatyna medac, 5 mg/ml, koncentrat do sporządzania roztworu do infuzji	DE/H/3915/001	22785	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	PL
Oxaliplatin "Ebewe" 5 mg/ml por oldatos infúzióhoz	DK/H/1774/001	OGYI-T-20312/01	EBEWE PHARMA	HU
Oxaliplatin "Ebewe" 5 mg/ml por oldatos infúzióhoz	DK/H/1774/001	OGYI-T-20312/02	EBEWE PHARMA	HU
Oxaliplatin "Ebewe" 5	DK/H/1774/001	OGYI-T-20312/03	EBEWE PHARMA	HU

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mg/ml por oldatos infúzióhoz				
Oxaliplatin "Ebewe", pulver til infusionsvæske, opløsning	DK/H/1774/001	39337	EBEWE PHARMA	DK
Oxaliplatin “Ebewe” 5 mg/ml powder for solution for infusion	AT/H/0185/001	20080014	EBEWE PHARMA	BG
Oxaliplatin ”Accord”, koncentrat til infusionsvæske, opløsning	UK/H/1349/001	44183	ACCORD HEALTHCARE LIMITED	DK
Oxaliplatin ”Accord”, koncentrat til infusionsvæske, opløsning	UK/H/1349/001	44183	ACCORD HEALTHCARE LIMITED	DK
Oxaliplatin ”Accord”, koncentrat til infusionsvæske, opløsning	UK/H/1349/001	44183	ACCORD HEALTHCARE LIMITED	DK
Oxaliplatin ”Actavis”, koncentrat til infusionsvæske, opløsning	UK/H/3700/001	46888	ACTAVIS GROUP PTC EHF.	DK
Oxaliplatin ”Actavis”, pulver til infusionsvæske, opløsning	DK/H/2505/001	40540	ACTAVIS GROUP HF.	DK
Oxaliplatin ”Fresenius Kabi”, koncentrat til infusionsvæske, opløsning	NL/H/4321/001	41945	FRESENIUS KABI ONCOLOGY PLC.	DK
Oxaliplatin ”Hospira”, koncentrat til infusionsvæske, opløsning	UK/H/0971/001	39930	HOSPIRA UK LTD	DK

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Oxaliplatin "Hospira", koncentrat til infusionsvæske, opløsning	UK/H/0971/001	39930	HOSPIRA UK LTD	DK
Oxaliplatin "Hospira", koncentrat til infusionsvæske, opløsning	UK/H/0971/001	39930	HOSPIRA UK LTD	DK
Oxaliplatin "Teva", koncentrat til infusionsvæske, opløsning	NL/H/0820/001	39379	TEVA DENMARK A/S	DK
Oxaliplatin 5 mg/ml concentrate for solution for infusion	NL/H/4321/001	PA 1422/004/001	FRESENIUS KABI ONCOLOGY PLC.	IE
Oxaliplatin 5 mg/ml concentrate for solution for infusion	UK/H/4547/001	PL 31750/0048	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	UK
Oxaliplatin 5 mg/ml concentrate for solution for infusion	-	PL 40378/0015	APTIL PHARMA LIMITED	UK
Oxaliplatin 5 mg/ml concentrate for solution for infusion	NL/H/4321/001	PL 18727/0016	FRESENIUS KABI ONCOLOGY PLC.	UK
Oxaliplatin 5 mg/ml Concentrate for Solution for Infusion	AT/H/0893/001	PL 18727/0012	FRESENIUS KABI ONCOLOGY PLC.	UK
Oxaliplatin 5 mg/ml powder for solution for infusion	AT/H/0185/001	MA586/00201	EBEWE PHARMA	MT
OXALIPLATIN 5 MG/ML POWDER FOR SOLUTION FOR INFUSION	FR/H/0144/001	PL 17780/0527	AVENTIS PHARMA LTD	UK
OXALIPLATIN 5 MG/ML POWDER FOR SOLUTION	FR/H/0144/001	PL 17780/0527	AVENTIS PHARMA LTD	UK

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FOR INFUSION				
OXALIPLATIN 5 MG/ML POWDER FOR SOLUTION FOR INFUSION	FR/H/0144/001	PL 17780/0527	AVENTIS PHARMA LTD	UK
Oxaliplatin 5 mg/ml, concentrate for solution for infusion	NL/H/0820/001	PL 00289/0310	TEVA UK LIMITED	UK
Oxaliplatin 5mg/ml Concentrate for Solution for Infusion	UK/H/1349/001	PA 1390/23/1	ACCORD HEALTHCARE LIMITED	IE
Oxaliplatin 5mg/ml Concentrate for Solution for Infusion	UK/H/1349/001	PA 1390/23/1	ACCORD HEALTHCARE LIMITED	IE
Oxaliplatin 5mg/ml Concentrate for Solution for Infusion	UK/H/1349/001	PA 1390/23/1	ACCORD HEALTHCARE LIMITED	IE
Oxaliplatin 5mg/ml concentrate for Solution for Infusion	UK/H/1349/001	MA 054/01701	ACCORD HEALTHCARE LIMITED	MT
Oxaliplatin 5mg/ml concentrate for Solution for Infusion	UK/H/1349/001	MA 054/01701	ACCORD HEALTHCARE LIMITED	MT
Oxaliplatin 5mg/ml concentrate for Solution for Infusion	UK/H/1349/001	MA 054/01701	ACCORD HEALTHCARE LIMITED	MT
Oxaliplatin 5mg/ml Concentrate for Solution for Infusion	UK/H/3700/001	PL 30306/0304	ACTAVIS GROUP PTC EHF.	UK
Oxaliplatin 5mg/ml concentrate for Solution for Infusion	UK/H/1349/001	PL 20075/0112	ACCORD HEALTHCARE LIMITED	UK
Oxaliplatin 5mg/ml concentrate for Solution for Infusion	UK/H/1349/001	PL 20075/0112	ACCORD HEALTHCARE LIMITED	UK
Oxaliplatin 5mg/ml	UK/H/1349/001	PL 20075/0112	ACCORD HEALTHCARE	UK

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concentrate for Solution for Infusion			LIMITED	
Oxaliplatin 5mg/ml powder for solution for infusion	NL/H/2337/001	PL 42069/0001	CADIASUN PHARMA GMBH	UK
Oxaliplatin Accord 5 mg/ml concentrat pentru solutie perfuzabila	UK/H/1349/001	2708/2010/01	ACCORD HEALTHCARE LIMITED	RO
Oxaliplatin Accord 5 mg/ml concentrat pentru soluţie perfuzabilă	UK/H/1349/001	2708/2010/02	ACCORD HEALTHCARE LIMITED	RO
Oxaliplatin Accord 5 mg/ml concentrat pentru soluţie perfuzabilă	UK/H/1349/001	2708/2010/03	ACCORD HEALTHCARE LIMITED	RO
Oxaliplatin Accord 5 mg/ml koncentrát pro prípravu infuzního roztoku	UK/H/1349/001	44/316/10-C	ACCORD HEALTHCARE LIMITED	CZ
Oxaliplatin Accord 5 mg/ml koncentrát pro prípravu infuzního roztoku	UK/H/1349/001	44/316/10-C	ACCORD HEALTHCARE LIMITED	CZ
Oxaliplatin Accord 5 mg/ml koncentrát pro prípravu infuzního roztoku	UK/H/1349/001	44/316/10-C	ACCORD HEALTHCARE LIMITED	CZ
Oxaliplatin Accord 5 mg/ml koncentrat till infusionsvätska, Lösning	UK/H/1349/001	41633	ACCORD HEALTHCARE LIMITED	SE
Oxaliplatin Accord 5 mg/ml koncentrat till infusionsvätska, Lösning	UK/H/1349/001	41633	ACCORD HEALTHCARE LIMITED	SE
Oxaliplatin Accord 5 mg/ml koncentrat till infusionsvätska, Lösning	UK/H/1349/001	41633	ACCORD HEALTHCARE LIMITED	SE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxaliplatin Accord 5 mg/ml koncentratas infuziniam tirpalui	UK/H/1349/001	LT/1/10/2144/001	ACCORD HEALTHCARE LIMITED	LT
Oxaliplatin Accord 5 mg/ml koncentratas infuziniam tirpalui	UK/H/1349/001	LT/1/10/2144/002	ACCORD HEALTHCARE LIMITED	LT
Oxaliplatin Accord 5 mg/ml koncentratas infuziniam tirpalui	UK/H/1349/001	LT/1/10/2144/003	ACCORD HEALTHCARE LIMITED	LT
Oxaliplatin Accord 5 mg/ml koncentrāts infūziju šķīduma pagatavošanai	UK/H/1349/001	10-0151	ACCORD HEALTHCARE LIMITED	LV
Oxaliplatin Accord 5 mg/ml koncentrāts infūziju šķīduma pagatavošanai	UK/H/1349/001	10-0151	ACCORD HEALTHCARE LIMITED	LV
Oxaliplatin Accord 5 mg/ml koncentrāts infūziju šķīduma pagatavošanai	UK/H/1349/001	10-0151	ACCORD HEALTHCARE LIMITED	LV
Oxaliplatin Accord 5 mg/ml koncentrátum oldatos infúzióhoz	UK/H/1349/001	OGYI-T-21479/03	ACCORD HEALTHCARE LIMITED	HU
Oxaliplatin Accord 5 mg/ml koncentrátum oldatos infúzióhoz	UK/H/1349/001	OGYI-T-21479/01	ACCORD HEALTHCARE LIMITED	HU
Oxaliplatin Accord 5 mg/ml koncentrátum oldatos infúzióhoz	UK/H/1349/001	OGYI-T-21479/02	ACCORD HEALTHCARE LIMITED	HU
Oxaliplatin Accord 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	UK/H/1349/001	1-29637	ACCORD HEALTHCARE LIMITED	AT
Oxaliplatin Accord 5	UK/H/1349/001	1-29637	ACCORD HEALTHCARE	AT

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mg/ml Konzentrat zur Herstellung einer Infusionslösung			LIMITED	
Oxaliplatin Accord 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	UK/H/1349/001	1-29637	ACCORD HEALTHCARE LIMITED	AT
Oxaliplatin Accord 5 mg/ml, infusioonilahuse kontsentraat	UK/H/1349/001	674810	ACCORD HEALTHCARE LIMITED	EE
Oxaliplatin Accord 5 mg/ml, infusioonilahuse kontsentraat	UK/H/1349/001	674810	ACCORD HEALTHCARE LIMITED	EE
Oxaliplatin Accord 5 mg/ml, infusioonilahuse kontsentraat	UK/H/1349/001	674810	ACCORD HEALTHCARE LIMITED	EE
Oxaliplatin Accord 5 mg/ml, infuusiokonsentraatti, liuosta varten.	UK/H/1349/001	26943	ACCORD HEALTHCARE LIMITED	FI
Oxaliplatin Accord 5 mg/ml, infuusiokonsentraatti, liuosta varten.	UK/H/1349/001	26943	ACCORD HEALTHCARE LIMITED	FI
Oxaliplatin Accord 5 mg/ml, infuusiokonsentraatti, liuosta varten.	UK/H/1349/001	26943	ACCORD HEALTHCARE LIMITED	FI
Oxaliplatin Accord 5 mg/ml, Konzentrat zur Herstellung einer Infusionslösung	UK/H/1349/001	76427.00.00	ACCORD HEALTHCARE LIMITED	DE
Oxaliplatin Accord 5 mg/ml, Konzentrat zur Herstellung einer	UK/H/1349/001	76427.00.00	ACCORD HEALTHCARE LIMITED	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Infusionslösung				
Oxaliplatin Accord 5 mg/ml, Konzentrat zur Herstellung einer Infusionslösung	UK/H/1349/001	76427.00.00	ACCORD HEALTHCARE LIMITED	DE
Oxaliplatin Accord Healthcare 5 mg/ml concentraat voor oplossing voor infusie	UK/H/1349/001	BE373992	ACCORD HEALTHCARE LIMITED	BE
Oxaliplatin Accord Healthcare 5 mg/ml concentraat voor oplossing voor infusie	UK/H/1349/001	BE374001	ACCORD HEALTHCARE LIMITED	BE
Oxaliplatin Accord Healthcare 5 mg/ml concentraat voor oplossing voor infusie	UK/H/1349/001	BE418555	ACCORD HEALTHCARE LIMITED	BE
Oxaliplatin Accord Healthcare 5 mg/ml solution à diluer pour perfusion	UK/H/1349/001	BE373992	ACCORD HEALTHCARE LIMITED	BE
Oxaliplatin Accord Healthcare 5 mg/ml solution à diluer pour perfusion	UK/H/1349/001	BE374001	ACCORD HEALTHCARE LIMITED	BE
Oxaliplatin Accord Healthcare 5 mg/ml solution à diluer pour perfusion	UK/H/1349/001	BE418555	ACCORD HEALTHCARE LIMITED	BE
Oxaliplatin Actavis 5 mg/ml concentrat pentru soluţie perfuzabilă	UK/H/3700/001	3641/2011/01-02-03	ACTAVIS GROUP PTC EHF.	RO
Oxaliplatin Actavis 5 mg/ml innrennslisþykkni, lausn	UK/H/3700/001	IS/1/11/065/01	ACTAVIS GROUP PTC EHF.	IS

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxaliplatin Actavis 5 mg/ml koncentratas infuziniam tirpalui	UK/H/3700/001	LT/1/08/0997/003-005	ACTAVIS GROUP PTC EHF.	LT
OXALIPLATIN ACTAVIS 5 mg/ml koncentrāts infūziju šķīduma pagatavošanai	UK/H/3700/001	11-0268	ACTAVIS GROUP PTC EHF.	LV
Oxaliplatin Actavis 5 mg/ml konsentrat til infusjonsvæske, oppløsning	UK/H/3700/001	10-7598	ACTAVIS GROUP PTC EHF.	NO
Oxaliplatin Actavis 5 mg/ml pulbere pentru soluţie perfuzabilă	not available	6169/2006/01-02	ACTAVIS SRL	RO
Oxaliplatin Actavis 5 mg/ml pulbere pentru soluţie perfuzabilă	not available	6169/2006/01-02	ACTAVIS SRL	RO
Oxaliplatin Actavis 5 mg/ml Pulver zur Herstellung einer Infusionslösung	DK/H/2505/001	1-27386	ACTAVIS GROUP HF.	AT
Oxaliplatin Actavis 5mg/ml koncentrátum oldatos infúzióhoz	UK/H/3700/001/DC	OGYI-T-21943/01-06	ACTAVIS GROUP PTC EHF.	HU
Oxaliplatin Actavis, 5 mg/ml infusioonilahuse kontsentraat	UK/H/3700/001	744711	ACTAVIS GROUP PTC EHF.	EE
Oxaliplatin AqVida 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	2200078.00.00	AQVIDA GMBH	DE
Oxaliplatin AqVida 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	2200079.00.00	AQVIDA GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxaliplatin AqVida 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	2200076.00.00	AQVIDA GMBH	DE
Oxaliplatin AqVida 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	2200077.00.00	AQVIDA GMBH	DE
Oxaliplatin AqVida 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	88845.00.00	AQVIDA GMBH	DE
Oxaliplatin Aurobindo 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	UK/H/3700/001	82014.00.00	PUREN PHARMA GMBH & CO. KG	DE
Oxaliplatin Bendalis 5 mg/ml, Konzentrat zur Herstellung einer Infusionslösung	not available	81696.00.00	BENDALIS GMBH	DE
Oxaliplatin beta 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	86708.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Oxaliplatin Boston Biopharma 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	2200080.00.00	BOSTON BIOPHARMA LTD.	DE
Oxaliplatin Ebewe 5 mg/ml Pulver zur Herstellung einer Infusionslösung	AT/H/0185/001	1-26435	EBEWE PHARMA	AT
Oxaliplatin Ebewe 5mg/ml concentrate for	AT/H/0341/001	20120240	EBEWE PHARMA	BG

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
solution infusion				
Oxaliplatin Fresenius Kabi 5 mg/ml konsentrat til infusjonsvæske, oppløsning	NL/H/4321/001	07-5183	FRESENIUS KABI ONCOLOGY PLC.	NO
Oxaliplatin Fresenius Kabi 5 mg/ml, koncentrat till infusionsvätska, lösning	AT/H/0893/001	44409	FRESENIUS KABI ONCOLOGY PLC.	SE
Oxaliplatin Hospira 5 mg/ml Concentraat voor oplossing voor infusie	UK/H/0971/001	RVG 34481	HOSPIRA BENELUX BVBA	NL
Oxaliplatin Hospira 5 mg/ml Concentraat voor oplossing voor infusie	UK/H/0971/001	RVG 34481	HOSPIRA BENELUX BVBA	NL
Oxaliplatin Hospira 5 mg/ml Concentraat voor oplossing voor infusie	UK/H/0971/001	RVG 34481	HOSPIRA BENELUX BVBA	NL
Oxaliplatin Hospira 5 mg/ml Concentrate for Solution for Infusion	UK/H/0971/001	PA 0437/056/002	HOSPIRA UK LTD	IE
Oxaliplatin Hospira 5 mg/ml Concentrate for Solution for Infusion	UK/H/0971/001	PA 0437/056/002	HOSPIRA UK LTD	IE
Oxaliplatin Hospira 5 mg/ml Concentrate for Solution for Infusion	UK/H/0971/001	PA 0437/056/002	HOSPIRA UK LTD	IE
Oxaliplatin Hospira 5 mg/ml Concentrate for Solution for Infusion	UK/H/0971/001	PL 04515/0215	HOSPIRA UK LTD	UK
Oxaliplatin Hospira 5 mg/ml Concentrate for Solution for Infusion	UK/H/0971/001	PL 04515/0215	HOSPIRA UK LTD	UK
Oxaliplatin Hospira 5 mg/ml Concentrate for	UK/H/0971/001	PL 04515/0215	HOSPIRA UK LTD	UK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Solution for Infusion				
Oxaliplatin Hospira 5 mg/ml koncentratas infuziniam tirpalui	UK/H/0971/001	LT/1/08/0962/001	HOSPIRA UK LTD	LT
Oxaliplatin Hospira 5 mg/ml koncentratas infuziniam tirpalui	UK/H/0971/001	LT/1/08/0962/002	HOSPIRA UK LTD	LT
Oxaliplatin Hospira 5 mg/ml koncentratas infuziniam tirpalui	UK/H/0971/001	LT/1/08/0962/003	HOSPIRA UK LTD	LT
Oxaliplatin Hospira 5 mg/ml koncentrāts infūziju šķīduma pagatavošanai	UK/H/0971/001	07-0326	HOSPIRA UK LTD	LV
Oxaliplatin Hospira 5 mg/ml koncentrāts infūziju šķīduma pagatavošanai	UK/H/0971/001	07-0326	HOSPIRA UK LTD	LV
Oxaliplatin Hospira 5 mg/ml koncentrāts infūziju šķīduma pagatavošanai	UK/H/0971/001	07-0326	HOSPIRA UK LTD	LV
Oxaliplatin Hospira 5 mg/ml, koncentrat till infusionsvätska, lösning	UK/H/0971/001	24413	HOSPIRA UK LTD	SE
Oxaliplatin Hospira 5 mg/ml, koncentrat till infusionsvätska, lösning	UK/H/0971/001	24413	HOSPIRA UK LTD	SE
Oxaliplatin Hospira 5 mg/ml, koncentrat till infusionsvätska, lösning	UK/H/0971/001	24413	HOSPIRA UK LTD	SE
OXALIPLATIN HOSPIRA 5mg/ml . Πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση	UK/H/0971/001	20672	HOSPIRA UK LTD	CY

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
OXALIPLATIN HOSPIRA 5mg/ml . Πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση	UK/H/0971/001	20672	HOSPIRA UK LTD	CY
OXALIPLATIN HOSPIRA 5mg/ml . Πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση	UK/H/0971/001	20672	HOSPIRA UK LTD	CY
Oxaliplatin Hospira, 5 mg/ml infusioonilahuse kontsentraat	UK/H/0971/001	563307	HOSPIRA UK LTD	EE
Oxaliplatin Hospira, 5 mg/ml infusioonilahuse kontsentraat	UK/H/0971/001	563307	HOSPIRA UK LTD	EE
Oxaliplatin Hospira, 5 mg/ml infusioonilahuse kontsentraat	UK/H/0971/001	563307	HOSPIRA UK LTD	EE
Oxaliplatin Kabi 5 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0893/001	10109/2017/01	FRESENIUS KABI ONCOLOGY PLC.	RO
Oxaliplatin Kabi 5 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0893/001	10109/2017/02	FRESENIUS KABI ONCOLOGY PLC.	RO
Oxaliplatin Kabi 5 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0893/001	10109/2017/05	FRESENIUS KABI ONCOLOGY PLC.	RO
Oxaliplatin Kabi 5 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0893/001	10109/2017/03	FRESENIUS KABI ONCOLOGY PLC.	RO
Oxaliplatin Kabi 5 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0893/001	10109/2017/04	FRESENIUS KABI ONCOLOGY PLC.	RO
Oxaliplatin Kabi 5 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0893/001	10109/2017/06	FRESENIUS KABI ONCOLOGY PLC.	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxaliplatin Kabi 5 mg/ml koncentrát pro infuzní roztok	NL/H/4321/001	44/624/09-C	FRESENIUS KABI ONCOLOGY PLC.	CZ
Oxaliplatin Kabi 5 mg/ml koncentratas infuziniam tirpalui	AT/H/0893/001	LT/1/10/1980/004	FRESENIUS KABI ONCOLOGY PLC.	LT
Oxaliplatin Kabi 5 mg/ml koncentratas infuziniam tirpalui	AT/H/0893/001	LT/1/10/1980/005	FRESENIUS KABI ONCOLOGY PLC.	LT
Oxaliplatin Kabi 5 mg/ml koncentratas infuziniam tirpalui	AT/H/0893/001	LT/1/10/1980/003	FRESENIUS KABI ONCOLOGY PLC.	LT
Oxaliplatin Kabi 5 mg/ml koncentrāts infūziju šķīduma pagatavošanai	AT/H/0893/001	11-0054	FRESENIUS KABI ONCOLOGY PLC.	LV
Oxaliplatin Kabi 5 mg/ml koncentrátum oldatos infúzióhoz	NL/H/4321/001	OGYI-T-21085/01	FRESENIUS KABI ONCOLOGY PLC.	HU
Oxaliplatin Kabi 5 mg/ml koncentrátum oldatos infúzióhoz	NL/H/4321/001	OGYI-T-21085/02	FRESENIUS KABI ONCOLOGY PLC.	HU
Oxaliplatin Kabi 5 mg/ml koncentrátum oldatos infúzióhoz	NL/H/4321/001	OGYI-T-21085/05	FRESENIUS KABI ONCOLOGY PLC.	HU
Oxaliplatin Kabi 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	AT/H/0893/001	1-30553	FRESENIUS KABI ONCOLOGY PLC.	AT
Oxaliplatin Kabi 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	NL/H/4321/001	70865.00.00	FRESENIUS KABI ONCOLOGY PLC.	DE
Oxaliplatin Kabi 5 mg/ml Konzentrat zur Herstellung einer	AT/H/0893/001	2011110055	FRESENIUS KABI ONCOLOGY PLC.	LU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Infusionslösung				
Oxaliplatin Kabi 5 mg/ml, infusioonilahuse kontsentraat	AT/H/0893/001	728111	FRESENIUS KABI ONCOLOGY PLC.	EE
Oxaliplatin Kabi 5 mg/ml, infúzny koncentrát	NL/H/4321/001	44/0590/09-S	FRESENIUS KABI ONCOLOGY PLC.	SK
Oxaliplatin Kabi, 5 mg/ml, koncentrat do sporządzania roztworu do infuzji	NL/H/4321/001	17086	FRESENIUS KABI ONCOLOGY PLC.	PL
Oxaliplatin Labosuan 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	89074.00.00	LABOSUAN I+D S.L.	DE
Oxaliplatin Labosuan 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	89074.00.00	LABOSUAN I+D S.L.	DE
Oxaliplatin medac 5 mg/ml concentrate for solution for infusion	DE/H/3915/001	PL 11587/0086	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	UK
Oxaliplatin medac 5 mg/ml infuusiokuiva-aine, liuosta varten	FI/H/0584/001	20370	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	FI
Oxaliplatin medac 5 mg/ml powder for solution for infusion	FI/H/0584/001	PA0623/008/001	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	IE
Oxaliplatin medac 5 mg/ml prášok na infúzny roztok	FI/H/0584/001	44/0524/06-S	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	SK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxaliplatin medac 5 mg/ml pulver till infusionsvätska, lösning	FI/H/0584/001	20370	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	FI
Oxaliplatin Mylan 5 mg/ml koncentrat ghal soluzzjoni ghall-infuzjoni	NL/H/1977/001	MA767/01601	MYLAN S.A.S	MT
Oxaliplatin Mylan 5 mg/ml koncentrat ghal soluzzjoni ghall-infuzjoni	NL/H/1977/002	MA767/01602	MYLAN S.A.S	MT
Oxaliplatin Mylan 5 mg/ml konċentrat għal soluzzjoni għall-infużjoni	NL/H/1977/003	MA767/01603	MYLAN S.A.S	MT
Oxaliplatin Mylan 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	NL/H/1977/001	1-31541	ARCANA ARZNEIMITTEL GMBH	AT
Oxaliplatin Mylan 5 mg/ml πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση	NL/H/1977/001	021824	MYLAN S.A.S	CY
Oxaliplatin OMNICARE 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	89696.00.00	OMNICARE PHARMA GMBH	DE
Oxaliplatin onkovis 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	88846.00.00	ONKOVIS GMBH	DE
Oxaliplatin ORCAzwei 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	89073.00.00	ORCAZWEI GMBH	DE
Oxaliplatin Pfizer 5 mg/ml innrennslisþykkni,	UK/H/0971/001	IS/1/06/056/01	PFIZER APS	IS

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
lausn				
Oxaliplatin Pfizer 5 mg/ml innrennslisþykkni, lausn	UK/H/0971/001	IS/1/06/056/01	PFIZER APS	IS
Oxaliplatin Pfizer 5 mg/ml innrennslisþykkni, lausn	UK/H/0971/001	IS/1/06/056/01	PFIZER APS	IS
Oxaliplatin Pfizer 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	UK/H/0971/001	1-27249	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
Oxaliplatin Pfizer 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	UK/H/0971/001	1-27249	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
Oxaliplatin Pfizer 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	UK/H/0971/001	1-27249	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
Oxaliplatin Pfizer 5 mg/ml, infuusiokonsentraatti, liuosta varten	UK/H/0971/001	22648	PFIZER PFE FINLAND OY	FI
Oxaliplatin Pfizer 5 mg/ml, infuusiokonsentraatti, liuosta varten	UK/H/0971/001	22648	PFIZER PFE FINLAND OY	FI
Oxaliplatin Pfizer 5 mg/ml, infuusiokonsentraatti, liuosta varten	UK/H/0971/001	22648	PFIZER PFE FINLAND OY	FI
Oxaliplatin Pfizer 5 mg/ml, koncentrat till infusionsvätska, lösning oxaliplatin	UK/H/0971/001	22648	PFIZER PFE FINLAND OY	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxaliplatin Pfizer 5 mg/ml, koncentrat till infusionsvätska, lösning oxaliplatin	UK/H/0971/001	22648	PFIZER PFE FINLAND OY	FI
Oxaliplatin Pfizer 5 mg/ml, koncentrat till infusionsvätska, lösning oxaliplatin	UK/H/0971/001	22648	PFIZER PFE FINLAND OY	FI
Oxaliplatin Pharma Resources 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	90005.00.00	PHARMA RESOURCES GMBH	DE
Oxaliplatin Pharma Resources 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	-	88714.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
Oxaliplatin Pharmagen 5 mg/ml koncentrát pro infuzní roztok	not available	44/421/14-C	PHARMAGEN CZ S.R.O.	CZ
Oxaliplatin Pliva 5 mg/ml koncentrat za otopinu za infuziju	not available	HR-H-505929304	PLIVA HRVATSKA D.O.O.	HR
Oxaliplatin Ribosepharm 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	DE/H/2769/001	81219.00.00	HIKMA PHARMA GMBH	DE
Oxaliplatin Ribosepharm 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	90006.00.00	PHARMA RESOURCES GMBH	DE
Oxaliplatin Sandoz 5 mg/ml konsentrat til infusjonsvæske,	AT/H/0341/001	09-6984	SANDOZ A/S	NO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
oppløsning				
Oxaliplatin STADA 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	88850.00.00	STADAPHARM GMBH	DE
Oxaliplatin STADA 5 mg/ml por oldatos infúzióhoz	FI/H/0589/001	OGYI-T-20773/02	STADA ARZNEIMITTEL AG	HU
Oxaliplatin STADA 5 mg/ml por oldatos infúzióhoz	FI/H/0589/001	OGYI-T-20773/03	STADA ARZNEIMITTEL AG	HU
Oxaliplatin STADA 5 mg/ml por oldatos infúzióhoz	FI/H/0589/001	OGYI-T-20773/01	STADA ARZNEIMITTEL AG	HU
Oxaliplatin STADA 5 mg/ml pulbere pentru soluţie perfuzabilă	FI/H/0589/001	4202/2012/03	STADA ARZNEIMITTEL AG	RO
Oxaliplatin STADA 5 mg/ml pulbere pentru soluţie perfuzabilă	FI/H/0589/001	4202/2012/01	STADA ARZNEIMITTEL AG	RO
Oxaliplatin STADA 5 mg/ml pulbere pentru soluţie perfuzabilă	FI/H/0589/001	4202/2012/02	STADA ARZNEIMITTEL AG	RO
Oxaliplatin SUN 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	UK/H/4547/001	82817.00.00	SUN PHARMACEUTICALS GERMANY GMBH	DE
Oxaliplatin Teva 5 mg/ml, koncentrat till infusionsvätska, lösning	NL/H/0820/001	23844	TEVA SWEDEN AB	SE
Oxaliplatin Teva, 5 mg/ml, koncentrat do sporządzania roztworu do infuzji	NL/H/0820/001	14292	TEVA PHARMACEUTICALS POLSKA SP. Z O.O.	PL
Oxaliplatin Tillomed 5	not available	90003.00.00	TILLOMED PHARMA GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
mg/ml Konzentrat zur Herstellung einer Infusionslösung				
Oxaliplatin Vianex 5 mg/ml pulver till infusionsvätska, lösning	SE/H/0960/001	23806	VIANEX S.A.	SE
Oxaliplatin Vipharm® 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	81697.00.00	VIPHARM GMBH	DE
Oxaliplatin Vitane 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	90004.00.00	VITANE PHARMA GMBH	DE
Oxaliplatin Winthrop 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	FR/H/0284/002	63654.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Oxaliplatin Winthrop 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	FR/H/0284/002	63654.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Oxaliplatin Winthrop 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	FR/H/0284/002	63654.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Oxaliplatin/Actavis 5 mg/ml πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση	UK/H/3700/001	022410	ACTAVIS GROUP PTC EHF.	CY
Oxaliplatin/Actavis 5 mg/ml πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση.	UK/H/3700/001	55938/13/19-06-2014	ACTAVIS GROUP PTC EHF.	GR
OXALIPLATIN/HOSPIRA 5	UK/H/0971/001	79689/08/12-01-2009	HOSPIRA UK LTD	GR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
mg/ml Πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση				
OXALIPLATIN/HOSPIRA 5 mg/ml Πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση	UK/H/0971/001	79689/08/12-01-2009	HOSPIRA UK LTD	GR
OXALIPLATIN/HOSPIRA 5 mg/ml Πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση	UK/H/0971/001	79689/08/12-01-2009	HOSPIRA UK LTD	GR
OXALIPLATIN/MEDICUS ®	not available	26229/14	MEDICUS A.E	GR
Oxaliplatin/Teva 5 mg/ml, πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση	NL/H/0820/001	55775/21-08-2015	TEVA PHARMA B.V.	GR
Oxaliplatina Accord 5 mg/ml Concentrado para solução para perfusão	UK/H/1349/001	5273735	ACCORD HEALTHCARE LIMITED	PT
Oxaliplatina Accord 5 mg/ml Concentrado para solução para perfusão	UK/H/1349/001	5273743	ACCORD HEALTHCARE LIMITED	PT
Oxaliplatina Accord 5 mg/ml Concentrado para solução para perfusão	UK/H/1349/001	5422431	ACCORD HEALTHCARE LIMITED	PT
Oxaliplatina Actavis 5 mg/ml prášok na infúzny roztok	DK/H/2505/001	44/0080/08-S	ACTAVIS GROUP HF.	SK
Oxaliplatina Aurovitas 5 mg/ml concentrado para solução para perfusão	UK/H/3700/001	5388160	AUROVITAS UNIPESSOAL, LDA.	PT
Oxaliplatina Aurovitas 5 mg/ml concentrado para solução para perfusão	UK/H/3700/001	5388178	AUROVITAS UNIPESSOAL, LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxaliplatina Aurovitas 5 mg/ml concentrado para solução para perfusão	UK/H/3700/001	5388202	AUROVITAS UNIPESSOAL, LDA.	PT
Oxaliplatina Aurovitas 5 mg/ml pó para solução para perfusão	PT/H/1702/001	5087655	AUROVITAS UNIPESSOAL, LDA.	PT
Oxaliplatina Aurovitas 5 mg/ml pó para solução para perfusão	PT/H/1702/001	5087663	AUROVITAS UNIPESSOAL, LDA.	PT
Oxaliplatina Hospira 5 mg/ml infúzny koncentrát	UK/H/0971/001	44/0459/07-S	HOSPIRA UK LTD	SK
Oxaliplatina Hospira 5 mg/ml infúzny koncentrát	UK/H/0971/001	44/0459/07-S	HOSPIRA UK LTD	SK
Oxaliplatina Hospira 5 mg/ml infúzny koncentrát	UK/H/0971/001	44/0459/07-S	HOSPIRA UK LTD	SK
Oxaliplatina Kabi 5 mg/ml concentrado para solução para perfusão	NL/H/4321/001	5201132	FRESENIUS KABI ONCOLOGY PLC.	PT
Oxaliplatina Kabi 5 mg/ml concentrado para solução para perfusão	NL/H/4321/001	5396171	FRESENIUS KABI ONCOLOGY PLC.	PT
Oxaliplatina Kabi 5 mg/ml concentrado para solução para perfusão	NL/H/4321/001	5201124	FRESENIUS KABI ONCOLOGY PLC.	PT
Oxaliplatina medac 5 mg/ml infúzny koncentrát	DE/H/3915/001	44/0268/14-S	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	SK
Oxaliplatina Medac 5 mg/ml koncentrát pro infuzní roztok	DE/H/3915/001	44/423/14-C	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	CZ

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxaliplatina Mylan 5 mg/ml concentrado para solução para perfusão	NL/H/1977/001	5643259	MYLAN, LDA	PT
Oxaliplatina Mylan 5 mg/ml concentrado para solução para perfusão	NL/H/1977/001	5643267	MYLAN, LDA	PT
Oxaliplatina Mylan 5 mg/ml concentrado para solução para perfusão	NL/H/1977/001	5643275	MYLAN, LDA	PT
Oxaliplatina Mylan 5 mg/ml concentrat pentru solutie perfuzabila	NL/H/1977/001	9540/2016/01	MYLAN S.A.S	RO
Oxaliplatina Mylan 5 mg/ml concentrat pentru solutie perfuzabila	NL/H/1977/001	9540/2016/02	MYLAN S.A.S	RO
Oxaliplatina Mylan 5 mg/ml concentrat pentru solutie perfuzabila	NL/H/1977/001	9540/2016/03	MYLAN S.A.S	RO
Oxaliplatina Mylan 5 mg/ml prášek pro infuzní roztok	FR/H/0324/001	44/646/07-C	MYLAN S.A.S	CZ
Oxaliplatina Mylan 5 mg/ml prášok na infúzny roztok	FR/H/0324/001	44/0345/07-S	MYLAN S.A.S	SK
Oxaliplatina Stada 5 mg/ml Pó para solução para perfusão	FI/H/0589/001	5184031	STADA LDA.	PT
Oxaliplatina Stada 5 mg/ml Pó para solução para perfusão	FI/H/0589/001	5115407	STADA LDA.	PT
Oxaliplatina Stada 5 mg/ml Pó para solução para perfusão	FI/H/0589/001	5115415	STADA LDA.	PT
Oxaliplatine Accord 5 mg/ml concentraat voor	UK/H/1349/001	RVG 103779	ACCORD HEALTHCARE LIMITED	NL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
oplossing voor infusie				
Oxaliplatine Accord 5 mg/ml concentraat voor oplossing voor infusie	UK/H/1349/001	RVG 103779	ACCORD HEALTHCARE LIMITED	NL
Oxaliplatine Accord 5 mg/ml concentraat voor oplossing voor infusie	UK/H/1349/001	RVG 103779	ACCORD HEALTHCARE LIMITED	NL
OXALIPLATINE ACCORD 5 mg/ml, solution à diluer pour perfusion	UK/H/1349/001	34009 576 842 1 1	ACCORD HEALTHCARE FRANCE SAS	FR
OXALIPLATINE ACCORD 5 mg/ml, solution à diluer pour perfusion	UK/H/1349/001	34009 579 546 4 2	ACCORD HEALTHCARE FRANCE SAS	FR
OXALIPLATINE ACCORD 5 mg/ml, solution à diluer pour perfusion	UK/H/1349/001	34009 576 841 5 0	ACCORD HEALTHCARE FRANCE SAS	FR
OXALIPLATINE ARROW 5 mg/ml, poudre pour solution pour perfusion	PT/H/1702/001	NL33901	ARROW GENERIQUES	FR
OXALIPLATINE ARROW 5 mg/ml, solution à diluer pour perfusion	not available	39965	ARROW GENERIQUES	FR
OXALIPLATINE ARROW 5 mg/ml, solution à diluer pour perfusion	not available	39965	ARROW GENERIQUES	FR
Oxaliplatine Aurobindo 5 mg/ml, concentraat voor oplossing voor infusie	UK/H/3700/001	RVG 107512	AUROBINDO PHARMA B.V.	NL
Oxaliplatine Cadiasun 5 mg/ml poeder voor oplossing voor infusie	NL/H/2337/001	RVG 110045	CADIASUN PHARMA GMBH	NL
Oxaliplatine Fresenius Kabi 5 mg/ml concentraat voor oplossing voor infusie	AT/H/0893/001	BE384632	FRESENIUS KABI ONCOLOGY PLC.	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxaliplatine Fresenius Kabi 5 mg/ml concentraat voor oplossing voor infusie	AT/H/0893/001	BE428391	FRESENIUS KABI ONCOLOGY PLC.	BE
Oxaliplatine Fresenius Kabi 5 mg/ml concentraat voor oplossing voor infusie	AT/H/0893/001	BE384623	FRESENIUS KABI ONCOLOGY PLC.	BE
Oxaliplatine Fresenius Kabi 5 mg/ml concentraat voor oplossing voor infusie	NL/H/4321/001	RVG 100834	FRESENIUS KABI ONCOLOGY PLC.	NL
Oxaliplatine Fresenius Kabi 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	AT/H/0893/001	BE384632	FRESENIUS KABI ONCOLOGY PLC.	BE
Oxaliplatine Fresenius Kabi 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	AT/H/0893/001	BE428391	FRESENIUS KABI ONCOLOGY PLC.	BE
Oxaliplatine Fresenius Kabi 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	AT/H/0893/001	BE384623	FRESENIUS KABI ONCOLOGY PLC.	BE
Oxaliplatine Fresenius Kabi 5 mg/ml solution à diluer pour perfusion	AT/H/0893/001	BE384632	FRESENIUS KABI ONCOLOGY PLC.	BE
Oxaliplatine Fresenius Kabi 5 mg/ml solution à diluer pour perfusion	AT/H/0893/001	BE428391	FRESENIUS KABI ONCOLOGY PLC.	BE
Oxaliplatine Fresenius Kabi 5 mg/ml solution à diluer pour perfusion	AT/H/0893/001	BE384623	FRESENIUS KABI ONCOLOGY PLC.	BE
OXALIPLATINE HOSPIRA 5 mg/ml, solution à	UK/H/0971/001	34009 572 480 8 6	HOSPIRA FRANCE	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
diluer pour perfusion				
OXALIPLATINE HOSPIRA 5 mg/ml, solution à diluer pour perfusion	UK/H/0971/001	34009 572 482 0 8	HOSPIRA FRANCE	FR
OXALIPLATINE HOSPIRA 5 mg/ml, solution à diluer pour perfusion	UK/H/0971/001	34009 574 402 4 4	HOSPIRA FRANCE	FR
OXALIPLATINE HOSPIRA 5 mg/ml, solution à diluer pour perfusion	UK/H/0971/001	34009 572 696 0 9	HOSPIRA FRANCE	FR
OXALIPLATINE HOSPIRA 5 mg/ml, solution à diluer pour perfusion	UK/H/0971/001	34009 572 697 7 7	HOSPIRA FRANCE	FR
OXALIPLATINE HOSPIRA 5 mg/ml, solution à diluer pour perfusion	UK/H/0971/001	34009 574 403 0 5	HOSPIRA FRANCE	FR
OXALIPLATINE KABI 5 mg/ml, solution à diluer pour perfusion	not available	34009 575 927 3 8	FRESENIUS KABI ONCOLOGY PLC.	FR
OXALIPLATINE KABI 5 mg/ml, solution à diluer pour perfusion	not available	34009 575 929 6 7	FRESENIUS KABI ONCOLOGY PLC.	FR
OXALIPLATINE KABI 5 mg/ml, solution à diluer pour perfusion	not available	34009 576 512 1 3	FRESENIUS KABI ONCOLOGY PLC.	FR
OXALIPLATINE KABI 5 mg/ml, solution à diluer pour perfusion	not available	34009 576 513 8 1	FRESENIUS KABI ONCOLOGY PLC.	FR
OXALIPLATINE KABI 5 mg/ml, solution à diluer pour perfusion	not available	34009 580 364 3 9	FRESENIUS KABI ONCOLOGY PLC.	FR
OXALIPLATINE KABI 5 mg/ml, solution à diluer pour perfusion	not available	34009 580 366 6 8	FRESENIUS KABI ONCOLOGY PLC.	FR
OXALIPLATINE MEDAC 5	DE/H/3915/001	34009 586 857 1 2	MEDAC GESELLSCHAFT FÜR	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
mg/ml, solution à diluer pour perfusion			KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	
OXALIPLATINE MEDAC 5 mg/ml, solution à diluer pour perfusion	DE/H/3915/001	34009 586 858 8 0	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	FR
OXALIPLATINE MEDAC 5 mg/ml, solution à diluer pour perfusion	DE/H/3915/001	34009 586 859 4 1	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	FR
Oxaliplatine Mylan 5 mg/ml concentraat voor oplossing voor infusie	NL/H/1977/001	BE468062	MYLAN BVBA/SPRL	BE
Oxaliplatine Mylan 5 mg/ml concentraat voor oplossing voor infusie	NL/H/1977/001	BE468071	MYLAN BVBA/SPRL	BE
Oxaliplatine Mylan 5 mg/ml concentraat voor oplossing voor infusie	NL/H/1977/001	BE468080	MYLAN BVBA/SPRL	BE
Oxaliplatine Mylan 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung Oxaliplatin	NL/H/1977/001	BE468062	MYLAN BVBA/SPRL	BE
Oxaliplatine Mylan 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung Oxaliplatin	NL/H/1977/001	BE468071	MYLAN BVBA/SPRL	BE
Oxaliplatine Mylan 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung Oxaliplatin	NL/H/1977/001	BE468080	MYLAN BVBA/SPRL	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxaliplatine Mylan 5 mg/ml solution à diluer pour perfusion	NL/H/1977/001	BE468062	MYLAN BVBA/SPRL	BE
Oxaliplatine Mylan 5 mg/ml solution à diluer pour perfusion	NL/H/1977/001	BE468071	MYLAN BVBA/SPRL	BE
Oxaliplatine Mylan 5 mg/ml solution à diluer pour perfusion	NL/H/1977/001	BE468080	MYLAN BVBA/SPRL	BE
Oxaliplatine Mylan 5 mg/ml solution à diluer pour perfusion	NL/H/1977/001	201600069	MYLAN BVBA/SPRL	LU
Oxaliplatine Mylan 5 mg/ml, concentraat voor oplossing voor infusie	NL/H/1977/001	RVG 107390	MYLAN B.V.	NL
OXALIPLATINE MYLAN 5 mg/ml, poudre pour solution pour perfusion	FR/H/0324/001	NL 32863	MYLAN S.A.S	FR
OXALIPLATINE MYLAN 5 mg/ml, solution a diluer pour perfusion	NL/H/1977/001	NL 39768	MYLAN S.A.S	FR
Oxaliplatine SUN 5 mg/ml, concentraat voor oplossing voor infusie	UK/H/4547/001	RVG 108042	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	NL
OXALIPLATINE SUN 5 mg/ml, solution à diluer pour perfusion	UK/H/4547/001	34009 582 161 2 1	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	FR
OXALIPLATINE SUN 5 mg/ml, solution à diluer pour perfusion	UK/H/4547/001	34009 582 158 1 0	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	FR
OXALIPLATINE SUN 5 mg/ml, solution à diluer pour perfusion	UK/H/4547/001	34009 582 156 9 8	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	FR
OXALIPLATINE SUN 5 mg/ml, solution à diluer	UK/H/4547/001	34009 582 159 8 8	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
pour perfusion				
OXALIPLATINE SUN 5 mg/ml, solution à diluer pour perfusion	UK/H/4547/001	34009 582 160 6 0	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	FR
OXALIPLATINE SUN 5 mg/ml, solution à diluer pour perfusion	UK/H/4547/001	34009 582 157 5 9	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	FR
Oxaliplatine Teva 5 mg/ml concentraat voor oplossing voor infusie	NL/H/0820/003	BE303834	TEVA PHARMA BELGIUM N.V./S.A	BE
Oxaliplatine Teva 5 mg/ml concentraat voor oplossing voor infusie	NL/H/0820/002	BE303825	TEVA PHARMA BELGIUM N.V./S.A	BE
Oxaliplatine Teva 5 mg/ml concentraat voor oplossing voor infusie	NL/H/0820/004	BE318832	TEVA PHARMA BELGIUM N.V./S.A	BE
Oxaliplatine Teva 5 mg/ml concentraat voor oplossing voor infusie	NL/H/0820/001	BE303816	TEVA PHARMA BELGIUM N.V./S.A	BE
Oxaliplatine Teva 5 mg/ml solution à diluer pour perfusion.	NL/H/0820/001	2007110014	TEVA PHARMA BELGIUM N.V./S.A	LU
OXALIPLATINE TEVA 5 mg/ml, solution à diluer pour perfusion	NL/H/0820/001	NL32938	TEVA SANTÉ	FR
OXALIPLATINE WINTHROP 5 MG/ML CONCENTRAAT, CONCENTRAAT VOOR OPLOSSING VOOR INFUSIE	FR/H/0284/002	RVG 33127	SANOFI-AVENTIS NETHERLANDS B.V.	NL
OXALIPLATINE WINTHROP 5 MG/ML CONCENTRAAT, CONCENTRAAT VOOR	FR/H/0284/002	RVG 33127	SANOFI-AVENTIS NETHERLANDS B.V.	NL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
OPLOSSING VOOR INFUSIE				
OXALIPLATINE WINTHROP 5 MG/ML CONCENTRAAT, CONCENTRAAT VOOR OPLOSSING VOOR INFUSIE	FR/H/0284/002	RVG 33127	SANOFI-AVENTIS NETHERLANDS B.V.	NL
OXALIPLATINE WINTHROP 5 MG/ML, SOLUTION À DILUER POUR PERFUSION	FR/H/0284/002	567 117-6	SANOFI-AVENTIS FRANCE	FR
OXALIPLATINE WINTHROP 5 MG/ML, SOLUTION À DILUER POUR PERFUSION	FR/H/0284/002	569 815-2	SANOFI-AVENTIS FRANCE	FR
OXALIPLATINE WINTHROP 5 MG/ML, SOLUTION À DILUER POUR PERFUSION	FR/H/0284/002	567 115-3	SANOFI-AVENTIS FRANCE	FR
OXALIPLATIN-EBEWE, 5 MG/ML, PROSZEK DO SPORZĄDZANIA ROZTWORU DO INFUZJI	DK/H/1774/001	12614	EBEWE PHARMA	PL
Oxaliplatin-GRY® 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	NL/H/0820/001	66264.00.00	TEVA GMBH	DE
Oxaliplatino 5 mg / ml, concentrato per soluzione per infusione	NL/H/0820/001	038107013	TEVA ITALIA S.R.L.	IT
Oxaliplatino Accord 5 mg/ml concentrado para solución para perfusión EFG	UK/H/1349/001	72386	ACCORD HEALTHCARE S.L.U.	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxaliplatino Accord 5 mg/ml concentrado para solución para perfusión EFG	UK/H/1349/001	72386	ACCORD HEALTHCARE S.L.U.	ES
Oxaliplatino Accord 5 mg/ml concentrado para solución para perfusión EFG	UK/H/1349/001	72386	ACCORD HEALTHCARE S.L.U.	ES
Oxaliplatino Accord 5 mg/ml concentrato per soluzione per infusione	UK/H/1349/001	041274010	ACCORD HEALTHCARE LIMITED	IT
Oxaliplatino Accord 5 mg/ml concentrato per soluzione per infusione	UK/H/1349/001	041274022	ACCORD HEALTHCARE LIMITED	IT
Oxaliplatino Accord 5 mg/ml concentrato per soluzione per infusione	UK/H/1349/001	041274034	ACCORD HEALTHCARE LIMITED	IT
Oxaliplatino Aurobindo 5 mg/ml concentrato per soluzione per infusione	UK/H/3700/001	039999014	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Oxaliplatino Aurobindo 5 mg/ml concentrato per soluzione per infusione.	UK/H/3700/001	039999026	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Oxaliplatino Aurobindo 5 mg/ml concentrato per soluzione per infusione	UK/H/3700/001	039999038	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Oxaliplatino Aurovitas 5 mg/ml concentrado para solución para perfusión EFG	UK/H/3700/001	74.655	AUROVITAS SPAIN,S.A.U.	ES
Oxaliplatino Cadiasun 5 mg/ml polvo para solución para perfusión EFG	NL/H/2337/001/DC	77233	CADIASUN PHARMA GMBH	ES
Oxaliplatino GP-Pharm 5	not available	72666	GP-PHARM S.A.	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
mg/ml polvo para solución para perfusión EFG				
Oxaliplatino Hospira 5 mg/ml concentrado para solución para perfusión EFG	UK/H/0971/001	69518	HOSPIRA UK LTD	ES
Oxaliplatino Hospira 5 mg/ml concentrado para solución para perfusión EFG	UK/H/0971/001	69518	HOSPIRA UK LTD	ES
Oxaliplatino Hospira 5 mg/ml concentrado para solución para perfusión EFG	UK/H/0971/001	69518	HOSPIRA UK LTD	ES
Oxaliplatino Kabi 5 mg/ml concentrado para solución para perfusión EFG	NL/H/4321/001	71.346	FRESENIUS KABI ONCOLOGY PLC.	ES
Oxaliplatino Kabi 5 mg/ml concentrato per soluzione per infusione	NL/H/4321/001	039170028	FRESENIUS KABI ONCOLOGY PLC.	IT
Oxaliplatino Kabi 5 mg/ml concentrato per soluzione per infusione	NL/H/4321/001	039170030	FRESENIUS KABI ONCOLOGY PLC.	IT
Oxaliplatino Kabi 5 mg/ml concentrato per soluzione per infusione	NL/H/4321/001	039170016	FRESENIUS KABI ONCOLOGY PLC.	IT
Oxaliplatino Pfizer 5 mg/ml Concentrato per soluzione per infusione	UK/H/0971/001	038094037	PFIZER ITALIA S.R.L.	IT
Oxaliplatino Pfizer 5 mg/ml Concentrato per soluzione per infusione	UK/H/0971/001	038094013	PFIZER ITALIA S.R.L.	IT
Oxaliplatino Pfizer 5	UK/H/0971/001	038094025	PFIZER ITALIA S.R.L.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
mg/ml Concentrato per soluzione per infusione				
Oxaliplatino Strides Arcolab International 5 mg/ml concentrato per soluzione per infusione	NL/H/1977/001	043321013	MYLAN S.P.A.	IT
Oxaliplatino Strides Arcolab International 5 mg/ml concentrato per soluzione per infusione	NL/H/1977/001	043321025	MYLAN S.P.A.	IT
Oxaliplatino Strides Arcolab International 5 mg/ml concentrato per soluzione per infusione	NL/H/1977/001	043321037	MYLAN S.P.A.	IT
Oxaliplatino SUN 5 mg/ml concentrado para solución para perfusión EFG	UK/H/4547/001	75278	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	ES
Oxaliplatino SUN 5 mg/ml concentrato per soluzione per infusione	UK/H/4547/001	041761038	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	IT
Oxaliplatino SUN 5 mg/ml concentrato per soluzione per infusione	UK/H/4547/001	041761053	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	IT
Oxaliplatino SUN 5 mg/ml concentrato per soluzione per infusione	UK/H/4547/001	041761026	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	IT
Oxaliplatino SUN 5 mg/ml concentrato per soluzione per infusione	UK/H/4547/001	041761040	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	IT
Oxaliplatino SUN 5 mg/ml concentrato per soluzione per infusione	UK/H/4547/001	041761014	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxaliplatino SUN 5 mg/ml concentrato per soluzione per infusione	UK/H/4547/001	041761065	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	IT
Oxaliplatino Teva 5 mg / ml, concentrato per soluzione per infusione	NL/H/0820/001	038107025	TEVA ITALIA S.R.L.	IT
Oxaliplatino Teva 5 mg / ml, concentrato per soluzione per infusione	NL/H/0820/001	038107076	TEVA ITALIA S.R.L.	IT
Oxaliplatino Teva 5 mg / ml, concentrato per soluzione per infusione.	NL/H/0820/001	038107037	TEVA ITALIA S.R.L.	IT
Oxaliplatino TEVA 5 mg/ml concentrado para solución para perfusión EFG	NL/H/0820/001	69.288	TEVA PHARMA S.L.U	ES
Oxaliplatin-Teva 5 mg/ml koncentratas infuziniam tirpalui	NL/H/0820/001	LT/1/07/0921/004	PHARMACHEMIE B.V	LT
Oxaliplatin-Teva 5 mg/ml koncentratas infuziniam tirpalui	NL/H/0820/001	LT/1/07/0921/005	PHARMACHEMIE B.V	LT
Oxaliplatin-Teva 5 mg/ml koncentratas infuziniam tirpalui	NL/H/0820/001	LT/1/07/0921/006	PHARMACHEMIE B.V	LT
Oxaliplatin-Teva 5 mg/ml koncentratas infuziniam tirpalui	NL/H/0820/001	LT/1/07/0921/007	PHARMACHEMIE B.V	LT
OXALIPLATIN-TEVA 5 mg/ml koncentrāts infūziju šķīduma pagatavošanai	NL/H/0820/001	07-0292	PHARMACHEMIE B.V	LV
Oxaliplatin-Teva 5 mg/ml koncentrátum oldatos infúzióhoz	NL/H/0820/001	OGYI-T-20505/02	TEVA GYÓGYSZERGYÁR ZRT	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxaliplatin-Teva 5 mg/ml koncentrátum oldatos infúzióhoz	NL/H/0820/001	OGYI-T-20505/01	TEVA GYÓGYSZERGYÁR ZRT	HU
Oxaliplatin-Teva 5 mg/ml koncentrátum oldatos infúzióhoz	NL/H/0820/001	OGYI-T-20505/04	TEVA GYÓGYSZERGYÁR ZRT	HU
Oxaliplatin-Teva 5 mg/ml koncentrátum oldatos infúzióhoz	NL/H/0820/001	OGYI-T-20505/03	TEVA GYÓGYSZERGYÁR ZRT	HU
Oxaliplatin-Teva 5 mg/ml, infusioonilahuse kontsentraat	NL/H/0820/001	553907	PHARMACHEMIE B.V	EE
Oxaliplatin-Teva 5 mg/ml, koncentrát pro infuzní roztok	NL/H/0820/001	44/591/07-C	TEVA PHARMACEUTICALS CR, S.R.O.	CZ
Oxaliplatinum Accord, 5 mg/ml, koncentrat do sporządzania roztworu do infuzji	UK/H/1349/001	17070	ACCORD HEALTHCARE LIMITED	PL
Oxaliplatinum Accord, 5 mg/ml, koncentrat do sporządzania roztworu do infuzji	UK/H/1349/001	17070	ACCORD HEALTHCARE LIMITED	PL
Oxaliplatinum Accord, 5 mg/ml, koncentrat do sporządzania roztworu do infuzji	UK/H/1349/001	17070	ACCORD HEALTHCARE LIMITED	PL
OXALIPROL® 5mg/ml πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση	not available	91329/11/ 23-01-2014	PHARMAZAC SA	GR
Oxalisan 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	86711.00.00	MEDICOPHARM AG	DE

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Oxalisin 5 mg/ml, concentraat voor oplossing voor infusie.	NL/H/0820/001	RVG 34033	PHARMACHEMIE B.V	NL
OXALIZOR® 5mg/ml πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση	not available	36849/05-11-2015	VOCATE ΦΑΡΜΑΚΕΥΤΙΚΗ ΑΕ	GR
OXALTINA® 5mg/ml κόνις για διάλυμα προς έγχυση	not available	24861/13/06-06-2014	PHARMATHEN HELLAS S.A.	GR
Oxamed 5 mg/ml infuusiokuiva-aine, liuosta varten	FI/H/0589/001	21549	STADA ARZNEIMITTEL AG	FI
Oxaplamyl 5 mg/ml innrennslisþykkni, lausn	NL/H/1977/001	IS/1/11/096/01	MYLAN HOSPITAL AS	IS
Oxaplamyl 5 mg/ml koncentrat till infusionsvätska, lösning	NL/H/1977/001	44216	MYLAN HOSPITAL AS	SE
Oxaplamyl 5 mg/ml konsentrat til infusjonsvæske, oppløsning	NL/H/1977/001	10-7505	MYLAN HOSPITAL AS	NO
Oxaplamyl 5mg/ml infuusiokonsentraatti, liuosta varten	NL/H/1977/001	28630	MYLAN HOSPITAL AS	FI
Oxaplamyl, koncentrat til infusionsvæske, opløsning	NL/H/1977/001	46680	MYLAN HOSPITAL AS	DK
OXAVIATIN® 5 mg/ml concentrate for solution for infusion	not available	AA721/00601	VIANEX S.A.	MT
OXAVIATIN® 5 mg/ml κόνις για διάλυμα προς έγχυση	SE/H/0960/001	ML21057	VIANEX S.A.	CY
OXAVIATIN® 5 mg/ml	SE/H/0960/001	26163	VIANEX S.A.	GR

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κόνις για διάλυμα προς έγχυση				
OXAVIATIN® 5mg/ml πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση	not available	29264/16/23-03-2017	VIANEX S.A.	GR
PLAXITIN, 5mg/ml, πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση	AT/H/0341/001	74198/18-10-2012	EBEWE PHARMA	GR
RECTOXAL® 5 mg/ml κόνις για διάλυμα προς έγχυση	EL/H/0123/001	57125/14-11-2012	ARITI SA	GR
Riboxatin 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	80931.00.00	HIKMA PHARMA GMBH	DE
Riboxatin 5 mg/ml Pulver zur Herstellung einer Infusionslösung	DE/H/1900/001	66243.00.00	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	DE
Sinoxal 100 mg prašak za otopinu za infuziju	not available	HR-H-640588353	ACTAVIS GROUP PTC EHF.	HR
SINOXAL 5 MG/ML INFÚZNY KONCENTRÁT	UK/H/3700/001	44/0018/12-S	ACTAVIS GROUP PTC EHF.	SK
Sinoxal 5 mg/ml koncentrat za raztopino za infundiranje	UK/H/3700/001	H/11/01425/002	ACTAVIS GROUP PTC EHF.	SI
Sinoxal 5 mg/ml koncentrat za raztopino za infundiranje	UK/H/3700/001	H/11/01425/003	ACTAVIS GROUP PTC EHF.	SI
Sinoxal 5 mg/ml koncentrat za raztopino za infundiranje	UK/H/3700/001	H/11/01425/001	ACTAVIS GROUP PTC EHF.	SI
Sinoxal 5 mg/ml por oldatos infúzióhoz	HU/H/0558/001	OGYI-T-20694/02	ACTAVIS GROUP HF.	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Sinoxal 50 mg prašak za otopinu za infuziju	not available	HR-H-640588353	ACTAVIS GROUP PTC EHF.	HR
Sinoxal 5mg/ml por oldatos infúzióhoz	HU/H/0558/001	OGYI-T-20694/01	ACTAVIS GROUP HF.	HU
VELMINOX®	not available	3197	VIOFAR LTD	GR
Οξαλιπλατίνη medac 5 mg/ml κόνις για διάλυμα προς έγχυση	FI/H/0584/001	20211	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	CY
ОКСАЛИПЛАТИН АКОРД 5 mg/ml концентрат за инфузионен разтвор	UK/H/1349/001	20120332	ACCORD HEALTHCARE LIMITED	BG
ОКСАЛИПЛАТИН АКОРД 5 mg/ml концентрат за инфузионен разтвор	UK/H/1349/001	20120332	ACCORD HEALTHCARE LIMITED	BG
ОКСАЛИПЛАТИН АКОРД 5 mg/ml концентрат за инфузионен разтвор	UK/H/1349/001	20120332	ACCORD HEALTHCARE LIMITED	BG
Оксалиплатин Актавис 5 mg/ml концентрат за инфузионен разтвор	UK/H/3700/001	20110406	ACTAVIS GROUP PTC EHF.	BG
Оксалиплатин Булгермед ВЕ 5 mg/ml концентрат за инфузионен разтвор	not available	20170376	BULGERMED VE	BG
Оксалиплатин Каби 5 mg/ml концентрат за инфузионен разтвор	AT/H/0893/001	20120487	FRESENIUS KABI ONCOLOGY PLC.	BG
Оксалиплатин Майлен 5 mg/ml концентрат за инфузионен разтвор	NL/H/1977/001	20120352	MYLAN S.A.S	BG
Оксалиплатин-Чайкафарма 100 mg прах за инфузионен разтвор	not available	II-14340	TCHAIKAPHARMA HIGH QUALITY MEDICINES, INC.	BG

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Оксалиплатин-Чайкафарма 50 mg прах за инфузионен разтвор	not available	II-14339	TCHAIKAPHARMA HIGH QUALITY MEDICINES, INC.	BG